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What is it like to have type-2 blindsight? Drawing inferences from residual function in type-1 blindsight.

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Abstract

Controversy surrounds the question of whether the experience sometimes elicited by visual stimuli in blindsight (type-2 blindsight) is visual in nature or whether it is some sort of non-visual experience. The suggestion that the experience is visual seems, at face value, to make sense. I argue here, however, that the residual abilities found in type-1 blindsight (blindsight in which stimuli elicit no conscious experience) are not aspects of normal vision with consciousness deleted, but are based fragments of visual processes that, in themselves, would not be intelligible as visual experiences. If type-2 blindsight is a conscious manifestation of this residual function then it is not obvious that type-2 blindsight would be phenomenally like vision.

Keywords

Vision, Neuropsychology, Consciousness, Blindsight, Phenomenology.

1. Introduction.

Blindsight was originally characterised as “visual capacity remaining after damage to striate cortex...” “even though the patient had no awareness of ‘seeing’ in blind portion of his field” (Weiskrantz, et al, 1974). It was subsequently discovered that patients with blindsight did, under some circumstances, report some kind of experience associated with stimuli presented in their regions of blindness. This is now known as type-2 blindsight (e.g. Weiskrantz, 1997). The nature of type-2 blindsight has been controversial to say the least. If the experience reported in type-2 blindsight is like that of normal vision (something that Weiskrantz doubts) then this implies that conscious visual experience can be elicited in the absence of primary visual cortex (see e.g. ffytche & Zeki, 2011). The positive reports of experience associated with visual stimuli in type-2 blindsight need not, however, be like normal visual experiences. In experiments exploring type-2 blindsight the subject is typically asked to make decision about some property of a visual stimulus followed by a ‘commentary’ response indicating whether he or she had any awareness associated that stimulus. A positive response in this context might simply indicate ‘feeling of knowing’ – that is, a sense that the preceding discrimination decision was something other than a guess. Some phenomenal experience beyond a feeling of knowing may, however, also occur. In this paper will concentrate on the nature of any such phenomenal experience that may occur in type-2 blindsight.

One approach to answering this question is simply to assert that any phenomenal experience elicited by a visual stimulus will be, by definition, a visual experience. Others argue that this is not necessarily the case. A recent exchange of papers by Morten Overgaard (e.g. Overgaard & Grünbaum, 2011) and Berit Brogaard (e.g. Brogaard, 2011) explored these issues in depth. The possibility that a given type of experience may have multiple causes has been known since Muller coined his Law of Specific Nerve Energies in 1835. It is less clear whether a given cause, in this case a visual stimulus, can give rise to different types of experience. As Overgaard is at pains to point out, it is dangerous to reply on introspection. The degree to which blindsight report awareness and the ways in which they describe this awareness appear to be highly variable (see e.g. Zeki & ffytche, 1998) and may be influenced by their interpretations of their own condition and of the experimenter’s expectations. It is also important to distinguish between patients who display blindsight and those who simply have severely impaired normal vision where stimuli that would be clearly seen by normal observers are, instead, near the threshold of vision. Azzopardi and Cowey (1997) used a signal detection theoretic approach to demonstrate that blindsight (specifically type-1 blindsight) was qualitatively distinct from near-threshold vision. If we accept these two concerns we might ask what we can learn by studying the abilities, rather than subjective reports, of patients who clearly display type-1 blindsight.

The case I want to make is that the residual visual abilities of blindsight patients are so different from normal vision that it is hard to imagine what it is like to experience them. I will argue that ways in which blindsight subjects succeed in colour discrimination, shape discrimination, motion discrimination or luminance discrimination tasks bears little resemblance to those that would be used in normal vision. If blindsight subjects have experiences elicited by visual stimuli we can only

call these experiences 'vision' by asserting that they are by definition vision, not because they are like visual experience.

2. Colour and brightness.

In 1999 Morland published a paper reporting experiments that explicitly tested whether the effects of visual stimuli presented in the blind field of blindsight patient GY were comparable with his experiences of the same stimuli in his sighted field (Morland et al, 1999). In one set of experiments GY was asked to adjust a stimulus presented in his blind field so that it matched a stimulus presented in his good field. Separate tests were made of his ability to adjust the luminance, colour (wavelength of light) and speed of motion of stimuli so that they matched between blind and seeing fields. GY was able to match both wavelength and speed between left (sighted for GY) and right (blind for GY) hemifields with almost the same accuracy as a normal observer. Although GY could match the luminance of a pair of stimuli both presented in his blind visual field, he was unable to match luminance between stimuli presented in the blind and sighted hemifields. Morland concludes "the luminance-modulated percept derived from the hemianopic field is not mapped to a perceptual dimension that can be compared with normal brightness perception. The two percepts seem to be unrelated and uncoupled" (p.1189). Should we conclude that, even if GY's residual blind-field brightness processing is not like his normal brightness perception, nevertheless his residual motion and colour processing is like normal vision?

Morland includes some of GY's introspections about the manner in which he conducted the wavelength matching experiment. GY says "I make the stimulus neither too red nor too green compared to the stimulus in the normal field" (p. 1189) but when asked if his blind-field percepts were "the same as normal red or green. He responded by saying 'Nothing is the same; I just know I can do this match'" (p. 1190). So, even though he succeeds at the matching task his introspection suggests that he is not comparing the same kind of experiences.

Morland was asking GY to compare the wavelengths of lights presented in his blind and sighted fields. When we perceive colour normally, however, our experience depends upon much more than the wavelength of light reaching our eyes. When we perceive the colours of objects our percept is of a property of those objects in the world not a percept of the wavelength distribution of light reaching our retina. In perceiving colour the visual system estimates the efficiency with which a material reflects lights of different wavelengths. Roughly speaking a good reflector of long wavelength light looks red whilst a good reflector of short wavelength light looks blue. The spectrum of light that reaches our eyes from an object is not, however, solely determined by the reflectance properties of the object, it is also dependent on the wavelength composition of the light illuminating the object. The visual system takes account of variations in the spectral composition of lights illuminating objects so that their perceived colour remains relatively unaffected by changes in illumination – the process of colour constancy (see e.g. Smithson, 2005). Colour constancy allows us to judge whether two objects seen under different illuminants are made of the same material. It is important to realise, however, that colour constancy not only provides us with this cognitive ability but that it also affects our experience of colour. This is beautifully illustrated in some visual illusions prepared by Dale Purves and Beau Lotto (Purves & Lotto, 2003). We see what appears to be a pair of multi-coloured

Rubik's cubes, one viewed through yellow cellophane and one view through blue cellophane. Some blue tiles and some yellow tiles can clearly be seen on both. What is remarkable is that the blue tiles on the cube seen through the yellow filter and the yellow tiles seen through the blue filter project identical lights to our eyes yet the colours we *experience* when looking at them are quite different (blue and yellow). Our experiences of these blue and yellow colours do not change even when we know that the lights reaching our eyes from the two tiles are identical.

The anatomy of colour vision shows a clear progression from ganglion cells in the retina where neural responses are determined primarily by the wavelength composition of light, through striate cortex where cells responding to wavelength contrast are found and extrastriate areas that appear to compute colour constancy. Cerebral achromatopsics, that is, patients with cortical colour blindness, lack these later extrastriate colour areas, do not experience colour and cannot make covert colour discriminations (Heywood, Kentridge & Cowey, 1998a). These patients do, however, see the borders between regions of different colour (Heywood, Kentridge & Cowey, 1998b) and make decision about the similarity of stimuli based on the chromatic contrasts they make with their backgrounds rather than their surface colour (Kentridge, Heywood & Cowey, 2004). We subsequently showed that the blindsight patient DB, with damage to striate cortex, does not even respond on the basis of chromatic contrast, instead he simply makes matches on the basis of the wavelengths of light reflected by the patches being compared (Kentridge, Heywood & Weiskrantz, 2007). Purely chromatic stimuli that we, or even a cerebral achromatopsic, would judge as being similar are judged as being different by a blindsight patient. So, although blindsight patients can make judgments about the wavelength composition of light they do not go on to process this information in a manner that might yield a normal colour experience. They are even further removed from doing so than a cortically colour blind patient.

3. Motion

What of motion perception? In Morland's experiments GY performed extremely well when matching the speed of motion between stimuli in his sighted and blind visual fields. This does not, however, imply that he extracts information about motion from a stimulus in the same fashion as a normal observer. One might, for example, infer that an object has moved by noting its location at one time, wait with one's eyes shut, open them and note the object's position again, and finally compare the two positions. If the positions differ then one can infer that the object has moved without seeing any motion. It is easy to see that one could additionally make inferences about the speed and direction of motion without seeing any motion. This is, of course, not the normal way of seeing motion. Azzopardi & Cowey (2001) explored the mechanisms through which blindsight subjects extract information about motion using stimuli known as random dot kinematograms. A random dot kinematogram stimulus typically consists of a large number of randomly positioned dots all of which move. The dots may all move in different directions but, on average, there is a predominant direction of motion. In addition, each dot only moves a small distance before disappearing and being replaced by another dot at a different random location. It is not possible to judge the overall direction of motion by tracking changes in the position of any individual dot; indeed, it is hard to even do this, given the short lifetime of individual dots. Normal observers effortlessly see the direction and speed of motion in random dot

kinematograms. Blindsight patients fail entirely. If the elements of the random dot kinematogram initially change in contrast whilst remaining stationary before the onset of motion then blindsight patients cannot even determine when motion starts, let alone discriminate its speed or direction. Recently Azzopardi and Hock (2011) showed that the responses of blindsight patients to motion are influenced by changes in the contrast polarity of stimuli. If we see a black bar against a grey background extend upwards we see motion in an upward direction whether the extension to the bar is black or white. A blindsight patient reports *downward* motion when the upward extension is white. This pattern of performance is consistent with a response to motion energy rather than to changes in the location of a feature of an object over time. Normal subjects' perception of motion in random dot kinematograms is thought to rely on motion energy cues. So blindsight subjects may possess functioning motion energy detectors, but it appears that they are not always able to apply them when they might be useful and do apply them to in situations where normal observers would not. There are, then, situations where normal observers see motion but where blindsight patients cannot see the onset of motion, cannot discriminate the speed or direction of motion or even judge the motion to be in the opposite direction of that seen by the normal visual system. Some of the multiple mechanisms we use in motion perception may survive in blindsight but they are not deployed in the same manner as they are in normal motion vision.

4. Shape.

Morland did not test GY's ability to compare characteristics of the spatial structure of stimuli between visual fields. There are, however, telling results reported in Weiskrantz's 1986 monograph on blindsight (Weiskrantz, 1986). Blindsight patients had been shown to be able to discriminate both the orientation of lines (horizontal vs. vertical) and even shape such as 'X' and 'O' in the earliest reports of the phenomenon (Weiskrantz et al, 1974). In the later monograph Weiskrantz reports, however, that, despite being able to discriminate 'X' from 'O', patient DB could not discriminate equilateral triangles with their apex pointing upwards (e.g. Δ) from those with the apex pointing downwards. The ability to discriminate 'X' from 'O' might tempt one to infer that blindsight subjects can discriminate shape, the latter result with triangles shows that this is not the case. Notice that equilateral triangle with the apices pointing upwards or downwards are constructed from exactly the same set of lines. Blindsight subjects might be able to discriminate orientation or curvature but they do not appear to be able to extract the relationships between the components of stimuli and combine these into representations of shape. Again, what might appear to be a residual ability that is like normal vision is nothing of the sort. We may ask ourselves what it would be like to see the orientation and curvature of features in a scene but not to see the objects to which these features belong. For me, at least, this is not something I can imagine.

5. Conclusion.

Type-1 blindsight is, then, not normal vision with the consciousness deleted. It is not even fragments of normal vision with the consciousness deleted. Blindsight patients clearly have many residual abilities, but none of these are complete aspects of normal vision. If type-2 blindsight is a conscious manifestation of these residual visual functions the experience elicited by visual stimuli is likely to be visually unintelligible.

The profound differences between residual visual abilities in blindsight and normal vision may well explain why, if these residual functions are the basis of type-2 blindsight, patients are unwilling to describe experiences elicited by visual stimuli as being visual experiences. They would be unlike vision in the sense that they would not be consistent with experiences elicited by the same stimuli in normal vision. Can a case nevertheless still be made for claiming that these experiences are visual? One might argue that as the neural responses to visual stimuli still occur within the visual system of blindsight patients then any associated experiences will be visual in nature. Even this argument is not watertight. In a series of remarkable studies Sur (see e.g. von Melchner et al, 2000) showed that if axons from one eye of a ferret are redirected so that visual signals are sent to auditory cortex then the ferret still classifies signals detected by that eye as visual despite the fact that the neural responses elicited occur in auditory cortex. The cortical area within which a neural response occurs need not necessarily determine the modality of associated experience. Of course in these experiments the surgery for re-wiring of connections was performed when the ferrets were one day old and so the internal organisation of auditory cortex may have been modified during development.

I would conclude that the case for assuming experience in type-2 blindsight is like vision is weak. It is all too easy to interpret type-1 blindsight as vision with the consciousness deleted but we now know that the successes of blindsight patients in a variety of visual discrimination tasks are not evidence for the survival of normal visual abilities. We cannot know directly what the experiences of blindsight patients in type-2 mode are like. It may be the case that their experience is like vision but they say that it is not. To draw the inference that experience in type-2 blindsight is visual despite patients' denial requires a much stronger argument than the assertion that experiences elicited by visual stimuli are necessarily visual.

References.

- Azzopardi, P. & Cowey, A. (1997). Is blindsight like normal, near-threshold vision? *Proceedings of the National Academy of Sciences USA*, 94, 14190-14194.
- Azzopardi, P. & Cowey, A. (2001). Motion discrimination in cortically blind patients. *Brain*, 124, 30-46.
- Azzopardi, P. & Hock, H.S. (2011). Illusory motion perception in blindsight. *Proceedings of the National Academy of Sciences USA*, 108, 876-881.
- Brogaard, B. (2011). Are there unconscious perceptual processes? *Consciousness and Cognition*, 20, 449-463.
- ffytche, D.H. & Zeki, S. (2011). The primary visual cortex, and feedback to it, are not necessary for conscious vision. *Brain*, 134, 247-257.
- Morland, A.B., Jones, S.R., Finlay, A.L., Deyzac, E., Le, S., & Kemp, S. (1999). Visual perception of motion, luminance and colour in a human hemianope. *Brain*, 122, 1183-1198.
- Heywood, C.A., Kentridge, R.W., & Cowey, A. (1998a). Cortical color blindness is not 'blindsight for color'. *Consciousness and Cognition* 7, 410-423.
- Heywood, C.A., Kentridge, R.W., & Cowey, A. (1998b). Form and motion from colour in cerebral achromatopsia. *Experimental Brain Research* 123, 145-153.
- Kentridge, R.W., Heywood, C.A., & Cowey, A. (2004). Chromatic edges, surfaces & constancies in cerebral achromatopsia. *Neuropsychologia* 42, 821-830.
- Kentridge, R.W., Heywood, C.A., & Weiskrantz, L. (2007). Colour-contrast processing in human striate cortex. *Proceedings of the National Academy of Sciences USA* 104, 15129-15131.
- Overgaard, M., & Grünbaum, T. (2011). Consciousness and modality: On the possible preserved visual consciousness in blindsight subjects. *Consciousness and Cognition*, 20, 1855-1859.
- Purves, D. & Lotto, B. (2003). *Why We See What We Do: a Wholly Empirical Theory of Vision*. Sunderland Massachusetts: Sinauer Associates.
- Smithson, H.E. (2005). Sensory, computational and cognitive components of human colour constancy. *Philosophical Transactions of the Royal Society: Biological Sciences*, 360, 1329-1346.
- Weiskrantz, L. (1997). *Consciousness lost and found*. Oxford: Oxford University Press.

Weiskrantz, L. (1986). *Blindsight. A case study and implications*. Oxford: Oxford University Press.

Weiskrantz, L., Warrington, E.K., Sanders, M.D., & Marshall, J. (1974). Visual capacity in the hemianopic field following a restricted occipital ablation. *Brain*, 97, 709-728.

von Melchner, L., Pallas, S.L., & Sur, M. (2000). Visual behaviour mediated by retinal projections directed to the auditory pathway. *Nature*, 404, 871-876.

Zeki, S. & ffytche, D.H. (1998). The Riddoch syndrome: insights into the neurobiology of conscious vision. *Brain*, 121, 25-45.